

Ref #	Hits	Search Query	DBs	Default Operator	Plurals	Time Stamp
L1	889	536/53	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/11/14 09:17
L2	180	I1 and glycolipid	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/11/14 09:44
L3	179	I2 and (separat\$ or extract\$ or isolat\$)	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/11/14 09:44
L4	118	I3 and membrane	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/11/14 10:03
L5	116	I4 and (water or chloroform or methanol or pyridine)	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/11/14 09:45
L6	1	I3 and (semipermeable ADJ membrane)	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/11/14 09:44
L7	5618	glycolipid	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/11/14 09:44
L8	4656	I7 and (separat\$ or extract\$ or isolat\$)	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/11/14 09:44
L9	30	I8 and (semipermeable ADJ membrane)	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/11/14 10:03
L10	29	I9 and (water or chloroform or methanol or pyridine)	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/11/14 10:08
L11	1261	I8 and dialys\$	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/11/14 10:02
L12	1157	I11 and membrane	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/11/14 10:03

L13	11	I11 and (semipermeable ADJ membrane)	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/11/14 10:09
L14	1105	I12 and (water or chloroform or methanol or pyridine)	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/11/14 10:08
L15	597	I11 and (isotonic or osmo\$)	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/11/14 10:10

CODEN: RIKAAN; ISSN: 0370-5633

DT Journal

LA Japanese

AB Changes of glycolipids in the brain after formalin fixation were examined. Quantity of lipids in the brain decreased rapidly after formalin fixation. Glycolipids decreased to 50% 24 h after fixation, and to 10% after 4 mo. after fixation. Fatty acid composition of glycolipids showed a change characterized by both a diminution of long-chain fatty acids (C:23-27) 4 mo after fixation, and its change was more markedly noted in normal fatty acids than hydroxy fatty acids.

=> dis hist

(FILE 'HOME' ENTERED AT 11:25:23 ON 14 NOV 2005)

FILE 'CAPLUS' ENTERED AT 11:25:33 ON 14 NOV 2005

L1 148 S ISHIKAWA TAKAHIRO/AU
L2 2 S L1 AND GLYCOLIPID
L3 889 S YAMAGUCHI AKIRA/AU
L4 3 S L3 AND GLYCOLIPID
L5 75 S SUZUKI KYOKO/AU
L6 0 S L5 AND (GLYCOLIPID(W) SEPARA?)
L7 4 S L5 AND GLYCOLIPID

Connecting via Winsock to STN

Welcome to STN International! Enter x:x

LOGINID: ssspta1623kxg

PASSWORD :

TERMINAL (ENTER 1, 2, 3, OR ?):2

NEWS 1 Web Page URLs for STN Seminar Schedule - N. America
NEWS 2 "Ask CAS" for self-help around the clock
NEWS 3 JUL 20 Powerful new interactive analysis and visualization software, STN AnaVist, now available
NEWS 4 AUG 11 STN AnaVist workshops to be held in North America
NEWS 5 AUG 30 CA/CAPLUS - Increased access to 19th century research documents
NEWS 6 AUG 30 CASREACT - Enhanced with displayable reaction conditions
NEWS 7 SEP 09 ACD predicted properties enhanced in REGISTRY/ZREGISTRY
NEWS 8 OCT 03 MATHDI removed from STN
NEWS 9 OCT 04 CA/CAPLUS-Canadian Intellectual Property Office (CIPO) added to core patent offices
NEWS 10 OCT 06 STN AnaVist workshops to be held in North America
NEWS 11 OCT 13 New CAS Information Use Policies Effective October 17, 2005
NEWS 12 OCT 17 STN(R) AnaVist(TM), Version 1.01, allows the export/download of CAPLUS documents for use in third-party analysis and visualization tools
NEWS 13 OCT 27 Free KWIC format extended in full-text databases
NEWS 14 OCT 27 DIOGENES content streamlined
NEWS 15 OCT 27 EPFULL enhanced with additional content

NEWS EXPRESS JUNE 13 CURRENT WINDOWS VERSION IS V8.0, CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP), AND CURRENT DISCOVER FILE IS DATED 13 JUNE 2005

NEWS EXPRESS JUNE 13 CURRENT WINDOWS VERSION IS V8.0, CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP), AND CURRENT DISCOVER FILE IS DATED 13 JUNE 2005

NEWS HOURS	STN Operating Hours Plus Help Desk Availability
NEWS INTER	General Internet Information
NEWS LOGIN	Welcome Banner and News Items
NEWS PHONE	Direct Dial and Telecommunication Network Access to STN
NEWS WWW	CAS World Wide Web Site (general information)

Enter NEWS followed by the item number or name to see news on that specific topic.

All use of STN is subject to the provisions of the STN Customer agreement. Please note that this agreement limits use to scientific research. Use for software development or design or implementation of commercial gateways or other similar uses is prohibited and may result in loss of user privileges and other penalties.

FILE 'HOME' ENTERED AT 11:25:23 ON 14 NOV 2005

=> file caplus
COST IN U.S. DOLLARS

FULL ESTIMATED COST

SINCE FILE ENTRY	TOTAL SESSION
0.21	0.21

FILE 'CAPLUS' ENTERED AT 11:25:33 ON 14 NOV 2005
USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.
PLEASE SEE "HELP USAGETERMS" FOR DETAILS.
COPYRIGHT (C) 2005 AMERICAN CHEMICAL SOCIETY (ACS)

Copyright of the articles to which records in this database refer is held by the publishers listed in the PUBLISHER (PB) field (available for records published or updated in Chemical Abstracts after December 26, 1996), unless otherwise indicated in the original publications. The CA Lexicon is the copyrighted intellectual property of the American Chemical Society and is provided to assist you in searching databases on STN. Any dissemination, distribution, copying, or storing of this information, without the prior written consent of CAS, is strictly prohibited.

FILE COVERS 1907 - 14 Nov 2005 VOL 143 ISS 21
FILE LAST UPDATED: 13 Nov 2005 (20051113/ED)

Effective October 17, 2005, revised CAS Information Use Policies apply. They are available for your review at:

<http://www.cas.org/infopolicy.html>

=> s Ishikawa Takahiro/AU
L1 148 ISHIKAWA TAKAHIRO/AU

=> s 11 and glycolipid
8850 GLYCOLIPID
12685 GLYCOLIPIDS
15800 GLYCOLIPID
(GLYCOLIPID OR GLYCOLIPIDS)
L2 2 L1 AND GLYCOLIPID

=> dis 12 1-2 bib abs

L2 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN
AN 2004:631770 CAPLUS
DN 141:156386
TI Manufacture of glycolipids from coffee beans, and functional foods containing them
IN Ishikawa, Takahiro; Yamaguchi, Akira
PA Brooks Holdings K. K., Japan; Glyco Lipid Laboratory K. K.
SO Jpn. Kokai Tokkyo Koho, 7 pp.
CODEN: JKXXAF

DT Patent
LA Japanese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 2004217606	A2	20040805	JP 2003-10180	20030117
PRAI	JP 2003-10180		20030117		

AB Glycolipids are manufactured by extraction from coffee beans using organic solvents. An EtOH extract of coffee bean powder was evaporated, mixed with H₂O, centrifuged, and dried to give white powder containing glycolipids.

L2 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN
AN 2003:335114 CAPLUS
DN 138:334037

TI Method for separating glycolipids with mixture solvent
IN Ishikawa, Takahiro; Yamaguchi, Akira; Suzuki, Kyoko; Katsuyama, Kayoko
PA Japan

SO PCT Int. Appl., 23 pp.
CODEN: PIXXD2

DT Patent
LA Japanese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2003035658	A1	20030501	WO 2001-JP11281	20011221
	W: AU, CA, CN, IN, KR, RU, US				
	RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR				

JP 2003129083	A2	20030508	JP 2001-321157	20011018
US 2005119475	A1	20050602	US 2004-825210	20040416
PRAI JP 2001-321157	A	20011018		
WO 2001-JP11281	A	20011221		

AB A method for separating glycolipids (especially, gangliosides) is provided, with which a large number of samples are conveniently and economically treated, and many types of glycolipids are recovered with high yield. The method comprises: (a) a step for performing the hydrolysis treatment of the extract obtained by extracting a biol. sample (e.g., animal/plant cell, tissue, microorganism) with a mixture liquid of nonpolar solvents (e.g., chloroform, pyridine) and polar solvents (e.g., water, methanol), and bringing the sample solution obtained into a contact with a solution having the osmotic pressure lower than the sample solution via a semipermeable membrane; and (b) a step for continuing the contact until the sample solution is separated into two or three layers, and isolating the intermediate layer and/or the lower layer.

RE.CNT 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> s Yamaguchi Akira/AU
L3 889 YAMAGUCHI AKIRA/AU

=> s 13 and glycolipid
8850 GLYCOLIPID
12685 GLYCOLIPIDS
15800 GLYCOLIPID
(GLYCOLIPID OR GLYCOLIPIDS)
L4 3 L3 AND GLYCOLIPID

=> dis 14 1-4 bib abs

L4 ANSWER 1 OF 3 CAPLUS COPYRIGHT 2005 ACS on STN
AN 2004:631770 CAPLUS
DN 141:156386
TI Manufacture of glycolipids from coffee beans, and functional foods containing them
IN Ishikawa, Takahiro; Yamaguchi, Akira
PA Brooks Holdings K. K., Japan; Glyco Lipid Laboratory K. K.
SO Jpn. Kokai Tokkyo Koho, 7 pp.
CODEN: JKXXAF

DT Patent
LA Japanese

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
-----	-----	-----	-----	-----
PI JP 2004217606	A2	20040805	JP 2003-10180	20030117
PRAI JP 2003-10180		20030117		

AB Glycolipids are manufactured by extraction from coffee beans using organic solvents. An EtOH extract of coffee bean powder was evaporated, mixed with H2O, centrifuged, and dried to give white powder containing glycolipids.

L4 ANSWER 2 OF 3 CAPLUS COPYRIGHT 2005 ACS on STN
AN 2003:335114 CAPLUS
DN 138:334037

TI Method for separating glycolipids with mixture solvent
IN Ishikawa, Takahiro; Yamaguchi, Akira; Suzuki, Kyoko; Katsuyama, Kayoko
PA Japan
SO PCT Int. Appl., 23 pp.
CODEN: PIXXD2

DT Patent
LA Japanese

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
-----	-----	-----	-----	-----
PI WO 2003035658	A1	20030501	WO 2001-JP11281	20011221
W: AU, CA, CN, IN, KR, RU, US				
RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL,				

PT, SE, TR

JP 2003129083	A2	20030508	JP 2001-321157	20011018
US 2005119475	A1	20050602	US 2004-825210	20040416
PRAI JP 2001-321157	A	20011018		
WO 2001-JP11281	A	20011221		

AB A method for separating glycolipids (especially, gangliosides) is provided, with which a large number of samples are conveniently and economically treated, and many types of glycolipids are recovered with high yield. The method comprises: (a) a step for performing the hydrolysis treatment of the extract obtained by extracting a biol. sample (e.g., animal/plant cell, tissue, microorganism) with a mixture liquid of nonpolar solvents (e.g., chloroform, pyridine) and polar solvents (e.g., water, methanol), and bringing the sample solution obtained into a contact with a solution having the osmotic pressure lower than the sample solution via a semipermeable membrane; and (b) a step for continuing the contact until the sample solution is separated into two or three layers, and isolating the intermediate layer and/or the lower layer.

RE.CNT 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2005 ACS on STN
AN 2003:273751 CAPLUS
DN 139:358457
TI Plasmid-based gene transfer ameliorates visceral storage in a mouse model of Sandhoff disease
AU Yamaguchi, Akira; Katsuyama, Kayoko; Suzuki, Kyoko; Kosaka, Kenji; Aoki, Ichiro; Yamanaka, Shoji
CS School of Medicine, Yokohama City University, Yokohama, 236-0004, Japan
SO Journal of Molecular Medicine (Heidelberg, Germany) (2003), 81(3), 185-193
CODEN: JMLME8; ISSN: 0946-2716
PB Springer-Verlag
DT Journal
LA English
AB Sandhoff disease is a severe neurodegenerative disorder with visceral involvement caused by mutations in the HEXB gene coding for the β subunit of the lysosomal hexosaminidases A and B. HEXB mutations result in the accumulation of undegraded substrates such as GM2 and GA2 in lysosomes. We evaluated the efficacy of cationic liposome-mediated plasmid gene therapy using the Sandhoff disease mouse, an animal model of a human lysosomal storage disease. The mice received a single i.v. injection of two plasmids, encoding the human α and β subunits of hexosaminidase cDNAs. As a result, 10-35% of normal levels of hexosaminidase expression, theor. therapeutic levels, were achieved in most visceral organs, but not in the brain, 3 days after injection with decreased levels by day 7. Histochem. staining confirmed widespread enzyme activity in visceral organs. Both GA2 and GM2 were reduced by almost 10% and 50%, resp., on day 3, and by 60% and 70% on day 7 compared with untreated age-matched Sandhoff disease mice. Consistent with the biochem. results, a reduction in GM2 was observed in liver cells histol. as well. These initial findings support further development of the plasmid gene therapy against lysosomal diseases with visceral pathol.

RE.CNT 48 THERE ARE 48 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> s Suzuki Kyoko/AU
L5 75 SUZUKI KYOKO/AU

=> s 15 and (glycolipid(w) separa?)
8850 GLYCOLIPID
12685 GLYCOLIPIDS
15800 GLYCOLIPID
(GLYCOLIPID OR GLYCOLIPIDS)
352140 SEPARA?
274231 SEP
12584 SEPS
285623 SEP
(SEP OR SEPS)
446569 SEPD

1 SEPDS
446570 SEPD
(SEPD OR SEPDS)
92051 SEPG
1 SEPGS
92052 SEPG
(SEPG OR SEPGS)
555979 SEPN
36038 SEPNS
574211 SEPN
(SEPN OR SEPNS)
1378550 SEPARA?
(SEPARA? OR SEP OR SEPD OR SEPG OR SEPN)
68 GLYCOLIPID (W) SEPARA?
L6 0 L5 AND (GLYCOLIPID (W) SEPARA?)

=> s 15 and glycolipid
8850 GLYCOLIPID
12685 GLYCOLIPIDS
15800 GLYCOLIPID
(GLYCOLIPID OR GLYCOLIPIDS)

L7 4 L5 AND GLYCOLIPID

=> dis 17 1-4 bib abs

L7 ANSWER 1 OF 4 CAPLUS COPYRIGHT 2005 ACS on STN
AN 2003:335114 CAPLUS
DN 138:334037
TI Method for separating glycolipids with mixture solvent
IN Ishikawa, Takahiro; Yamaguchi, Akira; Suzuki, Kyoko; Katsuyama, Kayoko

PA Japan
SO PCT Int. Appl., 23 pp.
CODEN: PIXXD2

DT Patent
LA Japanese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2003035658	A1	20030501	WO 2001-JP11281	20011221
	W: AU, CA, CN, IN, KR, RU, US RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR				
	JP 2003129083	A2	20030508	JP 2001-321157	20011018
	US 2005119475	A1	20050602	US 2004-825210	20040416

PRAI JP 2001-321157 A 20011018
WO 2001-JP11281 A 20011221

AB A method for separating glycolipids (especially, gangliosides) is provided, with which a large number of samples are conveniently and economically treated, and many types of glycolipids are recovered with high yield. The method comprises: (a) a step for performing the hydrolysis treatment of the extract obtained by extracting a biol. sample (e.g., animal/plant cell, tissue, microorganism) with a mixture liquid of nonpolar solvents (e.g., chloroform, pyridine) and polar solvents (e.g., water, methanol), and bringing the sample solution obtained into a contact with a solution having the osmotic pressure lower than the sample solution via a semipermeable membrane; and (b) a step for continuing the contact until the sample solution is separated into two or three layers, and isolating the intermediate layer and/or the lower layer.

RE.CNT 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 2 OF 4 CAPLUS COPYRIGHT 2005 ACS on STN
AN 2003:273751 CAPLUS
DN 139:358457
TI Plasmid-based gene transfer ameliorates visceral storage in a mouse model of Sandhoff disease
AU Yamaguchi, Akira; Katsuyama, Kayoko; Suzuki, Kyoko; Kosaka, Kenji; Aoki, Ichiro; Yamanaka, Shoji

CS School of Medicine, Yokohama City University, Yokohama, 236-0004, Japan
SO Journal of Molecular Medicine (Heidelberg, Germany) (2003), 81(3), 185-193
CODEN: JMLME8; ISSN: 0946-2716
PB Springer-Verlag
DT Journal
LA English
AB Sandhoff disease is a severe neurodegenerative disorder with visceral involvement caused by mutations in the HEXB gene coding for the β subunit of the lysosomal hexosaminidases A and B. HEXB mutations result in the accumulation of undegraded substrates such as GM2 and GA2 in lysosomes. We evaluated the efficacy of cationic liposome-mediated plasmid gene therapy using the Sandhoff disease mouse, an animal model of a human lysosomal storage disease. The mice received a single i.v. injection of two plasmids, encoding the human α and β subunits of hexosaminidase cDNAs. As a result, 10-35% of normal levels of hexosaminidase expression, theor. therapeutic levels, were achieved in most visceral organs, but not in the brain, 3 days after injection with decreased levels by day 7. Histochem. staining confirmed widespread enzyme activity in visceral organs. Both GA2 and GM2 were reduced by almost 10% and 50%, resp., on day 3, and by 60% and 70% on day 7 compared with untreated age-matched Sandhoff disease mice. Consistent with the biochem. results, a reduction in GM2 was observed in liver cells histol. as well. These initial findings support further development of the plasmid gene therapy against lysosomal diseases with visceral pathol.

RE.CNT 48 THERE ARE 48 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 3 OF 4 CAPLUS COPYRIGHT 2005 ACS on STN
AN 1991:119653 CAPLUS
DN 114:119653
TI Biochemical analysis of cerebral leukoencephalopathy, with special reference to Nasu-Hakola's disease and atypical leukodystrophy
AU Suzuki, Kyoko
CS Sch. Med., Yokohama City Univ., Yokohama, 232, Japan
SO Yokohama Igaku (1990), 41(2), 163-72
CODEN: YKIGAK; ISSN: 0372-7726
DT Journal
LA Japanese
AB Changes of glycolipids were studied in 10 cases of cerebral leukoencephalopathy mainly consisting of Nasu-Hakola diseases and atypical leukodystrophies. In the demyelinated lesions, change was noticed in lipid, protein, and ganglioside, and fatty acid composition of glycolipids. Four stages were noticed in changes of the glycolipids in the demyelinated cerebral white matter. Stage 1: no change was noticed in the component of the glycolipids (norm. cerebroside:hydroxycerebroside:sulfatide, 1:1:1), in spite of a decrease of the total lipid content. Stage 2: decrease in the norm. cerebroside content was noticed (0.5:1:1). Stage 3: decrease in both norm. cerebroside and sulfatide (0.5:1:0.2) was noticed. Stage 4: hydroxycerebroside content was decreased resulting in complete loss of the glycolipid (0.1:0.1:trace). The content of the long chain fatty acids was significantly decreased in the cerebral cortex in the patients with leukoencephalopathies. Myelin degeneration in the cerebral white matter was divided into 2 types. In myelin-palor type, changes of the composition of the fatty acid were slight in spite of a marked decrease of the total glycolipid content in stage 4. Marked fibrillary gliosis was noted in the white matter in this type. In myelin-clastic type, very long chain fatty acid content was significantly decreased. Less fibrillary gliosis was seen in such a type.

L7 ANSWER 4 OF 4 CAPLUS COPYRIGHT 2005 ACS on STN
AN 1991:38619 CAPLUS
DN 114:38619
TI Changes of glycolipids in the human brain after formalin fixation
AU Suzuki, Kyoko; Yokoi, Susumu; Yamada, Yoshiteru; Arai, Nobutaka; Matsushita, Masaaki
CS Sch. Med., Yokohama City Univ., Yokohama, 236, Japan
SO Rinsho Kagaku (Nippon Rinsho Kagakkai) (1990), 19(2), 131-5